A Users’ Guide to the 2016 Surviving Sepsis Guidelines

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The 2016 Surviving Sepsis Guidelines have arrived, a remarkable document, all 67 pages with 655 references (1, 2). We congratulate the lead authors and contributing committee members. With each iteration, the guidelines grow more complex and perhaps more challenging to utilize. Herein, we offer guidance toward effective utilization.

LAYERS OF THE GUIDELINES

The guidelines may be thought of as several concentric layers, similar to an onion (Fig. 1).

The outer layer represents the recommendations. A bedside practitioner responsible for immediate decision making and trusting guidelines process will focus on the recommendations. This group of users may find the tables of abbreviated recommendations—the essence of the guidelines condensed to 7 pages—especially useful.

The next layer represents the rationales for the recommendations, illuminating the logic—the evidence and the thought—underlying each recommendation. For those who want a more in-depth understanding of how the recommendations were built, the rationales are a great resource. Moreover, the rationales help cement the recommendations for the busy practitioner: insight into the biologic plausibility and reasoning enable timely recall. The rationales also represent a foundation for educating healthcare practitioners on the recognition and treatment of sepsis.

The deepest layer, the core of the onion, houses the evidentiary tables. The tables compile and organize the existing data in a manner that provides insight into the reasoning behind each recommendation (magnitude of benefit or harm and the quality of evidence). This layer is typically for the inquisitive clinician and for the clinical scientist with focused interest in sepsis.

GUIDELINES AS A RESOURCE

The collected guidelines are a resource document applicable to a variety of areas of sepsis management. Some areas are broad, such as initial resuscitation. Some areas are narrow, such as empiric therapy of a potential fungal infection. Inspection and reflection will provide insight into what can be stated with confidence and—equally important—where opportunities for future research lie.

The guidelines also tell a story about the approach to treating the sepsis patient through a management continuum beginning with diagnosis, initial resuscitation, antimicrobial therapy, source control, fluid/vasoactive therapy, and progressing through organ support and adjunctive therapy recommendations.

Two aspects of the guidelines should be understood. We illuminate these two aspects through an analysis of the priority currently assigned to early identification and initial treatment of sepsis, including antibiotics and fluid therapy.

First, the recommendation for antibiotic administration within an hour of diagnosis of sepsis is a lofty goal of care, judged to be ideal for the patient but not yet standard care. Despite the best intentions of the healthcare team, antibiotic administration within one hour from time of diagnosis may be difficult due to the complexity of the hospital environment and essential care being delivered to other patients during the same time period by the same healthcare practitioners and health system. This is one among several “aspirational recommendations” considered by the experts to represent best practice that individual practitioners and healthcare teams should strive to operationalize.

Second, the clinician may push back from use of recommendations for fear that evidence-based guidelines lead to “cookie cutter” medicine and reflexive behaviors that deemphasize the “art” of medicine. The recommendations are intended for a “typical” septic patient. Patients still benefit from the art of medicine, which includes interpretation of data and individualization of treatment. The recommendations provide much-needed general treatment guidance to the bedside decision maker who is busy, pressured to see more patients in less time, and who will use a distillation of the current literature into a coherent set of recommendations suitable for the large majority of septic patients who are “typical”. For most of us in the
Foreword

Layers of the SSC Guidelines


Figure 1. The layers of an onion are paralleled to the components of the guidelines document, reflecting the depth of exploration by the user.

Figure 2. This figure explores the nuances of initial administration of 30 mL/kg crystalloid for sepsis-induced hypoperfusion based on patient characteristics. It also draws attention to reassessment tools following the initial fluid dose as an influence on further fluid administration or inotropic therapy.

trenches of everyday care, the lists of specific recommendations (seen in the tables in the manuscript) are a welcome adjunct to personalizing care.

This guidance includes sepsis management in the emergency department, the general hospital floors, and the ICU. For example, the recommendation for an initial 30 mL/kg crystalloid infusion for tissue hypoperfusion is chosen as a one value fit for bedside guidance. Administering 30 mL/kg crystalloid is a useful initial therapy for the majority of patients and this literature supported fluid dose is linked to good outcomes (3, 4). Figure 2 offers guidance for initial fluid

Application of Fluid Resuscitation in Adult Septic Shock

Sepsis-induced hypotension or lactate ≥ 4 mmol/L
(Based on SSC bundle and CMS threshold)

No high flow oxygen and
No ESRD on dialysis or CHF

Rapid Infusion
of 30 mL/kg
Crystalloid*

Pneumonia or ALI with high flow oxygen requirements

Intubated/mechanically ventilated

Consider intubation/mechanical ventilation to facilitate 30 mL/kg crystalloid *

Intubated/mechanically ventilated

Total of 30 mL/kg crystalloid* with frequent reassessment of oxygenation

Total of 30 mL/kg with frequent reassessment of oxygenation

If no

Rapid infusion of 30 mL/kg crystalloid *

Considerations post 30 mL/kg crystalloid infusion
1. Continue to balance fluid resuscitation and vasopressor dose with attention to maintain tissue perfusion and minimize interstitial edema
2. Implement some combination of the list below to aid in further resuscitation choices that may include additional fluid or inotrope therapy
   - blood pressure/heart rate response,
   - urine output,
   - cardiothoracic ultrasound,
   - CVP, ScvO2,
   - pulse pressure variation
   - lactate clearance/normalization or
   - dynamic measurement such as response of flow to fluid bolus or passive leg raising
3. Consider albumin fluid resuscitation, when large volumes of crystalloid are required to maintain intravascular volume.

*Administer 30 mL/kg crystalloid within first 3 hours

ALI=acute lung injury; CHF=congestive heart failure; CMS=US Centers for Medicare and Medicaid Services; CVP=central venous pressure; ESRD=end stage renal disease; kg=kilograms; ml=milliliters; oxyHb=oxyhemoglobin; ScvO2=superior vena cava oxygen saturation
resuscitation and is built forward from the guidelines recommendation for 30 mL/kg initial crystalloid fluid administration within the first six hours for sepsis-induced tissue hypoperfusion. The flow diagram incorporates some of our own opinions for successful fluid resuscitation based on experience and our understanding of the literature.

Another illustration is the recommendation for an initial mean arterial pressure target of septic shock of 65 mm Hg—a solid initial target with significant literature support—yet clearly one size does not fit all. Having a mean blood pressure target for the “typical” patient enables the art of medicine and provides a rationale for the provider in choosing a higher target for the atypical patient. Thus higher-than-reference values could—and perhaps even should—be selected for the patient with chronic poorly controlled hypertension, intra-abdominal compartment syndrome, or high central venous pressure (CVP) with acute decrease in renal perfusion (5–7).

**VALUES OF THE RECOMMENDATIONS**

What about strong versus weak recommendations? Strong recommendations should be included as part of usual care of the septic patient. Weak recommendations imply that although the majority of well-informed patients or surrogate decision makers would want this done, others would not. Recognizing the complexity of many septic patients—heterogeneity of disease process and co-morbidities—one may arrive at the conclusion that in a particular patient, even a strong recommendation may not be in that patient’s best interest.

What does the quality of evidence communicate that the strength of recommendation does not? The quality of evidence reflects the experts’ confidence in the recommendation: high quality evidence generally means that the experts have high confidence in the recommendation while low quality evidence reflects lower confidence in the recommendation. The quality of evidence is an important determinant of the strength of recommendation (“strong, do it” or “weak, probably do it”
recommendation). Substantial insight may be offered by the quality of evidence for the scientist searching for more information as to how he or she will use the recommendation to generate hypotheses for research.

How should a clinician use the best practice statement (BPS) recommendations? These are strong recommendations that lack evidence-based literature that likely will never be available because they are common sense—generally accepted good things to do for septic patients. For example, recommending that sepsis and septic shock treatment and resuscitation should begin immediately is common-sense good practice, and the alternative is implausible. BPS recommendations are also typically very low risk. BPS recommendations are formulated based on strict criteria, therefore, should be considered at least as strong as the strong recommendations.

Recommendations for resuscitation targets have gone from clear but controversial in 2012 to nuanced in the 2016 guidelines. Gone in 2016 are the specific targets of CVP and ScvO₂ to determine success of resuscitation, replaced with more general guidance as to a variety of targets (with emphasis on dynamic targets) that can be used. This is appropriate as it reflects the current lack of evidence as to a preferred target or approach to hemodynamic monitoring that deliver better clinical outcomes in sepsis (3, 4, 8). Because a preferred target is not known, a variety of reassessment options (after 30 mL/kg crystalloid fluid administration) should be considered.

APPLICATION OF RECOMMENDATIONS

All guidelines lead to questions. Here are a couple of common ones, and our personal approaches.

Question 1: “It is pretty clear that I should start out using norepinephrine as my initial vasopressor in septic shock—but where do I go from there using the other vasopressor recommendations?” Figure 3 offers guidance in this area and is constructed in compliance with the guidelines vasopressor recommendations.

Question 2: “When is my patient considered “hemodynamically unstable” after fluid administration and vasopressor initiation, as to warrant steroid administration?” A useful parallel here is the use of an inhaled selective pulmonary vasodilator in the severest of acute respiratory distress syndrome patients. This therapy improves oxygenation but does not improve outcome in multiple large randomized trials (9, 10). The same is true for trials of steroids for septic shock, which despite producing improvement in hemodynamics have no consistent positive effect on patient-important outcome (11, 12). So, consider these two low-risk therapies if there is concern that the patient will die of hypoxemia (acute respiratory distress syndrome) or hemodynamic instability (septic shock). Figure 3 incorporates steroid administration guidance into a vasopressor in septic shock flow diagram.

In closing, it is important to remember that the guidelines can be many things to many different user groups. As guidance
for the variety of users of the guidelines we offer Figure 4 as an approach to uncover the onion.

Enjoy your guidelines adventure!

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Note: Although the authors of this manuscript are members of the guidelines committee, the views expressed in this manuscript are from a personal perspective and do not represent any collective viewpoint of the guidelines committee.

REFERENCES